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(54) Title: A PROCESS FOR DEPOSITING A SiO₂ FILM HAVING REDUCED INTRINSIC STRESS AND/OR REDUCED HYDROGEN CONTENT

(57) Abstract

A process for reducing intrinsic stress and/or hydrogen content of a SiO₂ film grown by chemical vapor deposition. The process is applicable to plasma-enhanced and electron cyclotron resonance chemical vapor deposition of silicon dioxide wherein a vapor phase etchant is introduced while growing the silicon dioxide film. The presence of the etchant during the plasma deposition process allows for selective removal of high energy silicon dioxide molecules in the growing film thus reducing intrinsic stress within the film. The use of halogen etchants further reduces the amount of hydrogen present as hydroxyl within the film.

has stress
& halogensCF₄, C₂F₆, BF₃, Cl₂ etc (p.9 p.15)

PECVD (p.10, 11)

p.10 - control ratios of gases - p.15
p.13 TEOSlacks
dual frequency
conduction linesp.10 also equates use
of plasma to thermal
processes

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A PROCESS FOR DEPOSITING A SiO_x FILM HAVING REDUCED
INTRINSIC STRESS AND/OR REDUCED HYDROGEN CONTENT

BACKGROUND OF THE INVENTION

Field of the Invention

5 The present invention relates generally to
plasma-enhanced chemical vapor deposition (PECVD)
processes, and more particularly to a PECVD process for
growing silicon dioxide on a substrate surface wherein
the surface is simultaneously etched such that the
10 grown silicon dioxide layer has a reduced level of
intrinsic stress and/or a reduced hydrogen content.

Description of Related Technology

Chemical vapor deposition (CVD) is a process for
forming a material layer by the reaction of gas phase
15 reactants at or near a substrate surface. Thermal CVD
processes typically rely on heating of the surface in
order to promote the reaction(s) which result in
formation of compound(s) on the substrate surface. A
problem with thermal CVD processes is that the thermal
20 reaction occurs at high temperatures thus requiring
heat resistant substrates and it is difficult to
control characteristics of the film such as internal
stress. Depending on the nature of the deposited
material, the surface temperature required for
25 deposition may be as high as about 1000°C. Under a
variety of circumstances, it would be desirable to be

able to perform such CVD processes at lower temperatures.

In order to reduce the required surface temperatures, a variation of CVD referred to as plasma-enhanced CVD (PECVD) is employed. In PECVD, a radio frequency (RF) discharge converts the gas phase reactants to more reactive free radical species resulting in a plasma. The higher reactivity of the free radical species reduces the energy required for reaction and thus lowers the substrate surface temperature which is necessary to carry out the process. In particular, the specimen chamber includes a radio frequency electrode arranged opposite to the specimen table. Gas is introduced into the specimen chamber which is typically at a pressure of 0.1 to 10 Torr and radio frequency power is supplied to the specimen chamber to produce plasma. To grow a film of silicon nitride, gaseous molecules of SiH_4 and NH_3 can be dissociated into plasma and silicon nitride can be grown on a surface of a specimen substrate on a specimen table heated to 300-500°C. However, dissociation of SiH_4 and NH_3 may not be sufficient, H may be incorporated into the silicon nitride film and/or the Si-N bond may not be sufficient.

In a particular form of PECVD, referred to as electron cyclotron resonance (ECR) CVD, the plasma is established in a separate plasma chamber by the application of microwave energy to reactant molecules maintained within a magnetic field. In an ECR CVD apparatus, plasma is generated by a microwave discharge through electron cyclotron resonance and a thin film can be formed on a specimen surface with the

aid of a divergent magnetic field. In this case, molecules in the plasma are activated and reacted on a specimen substrate to form the thin film. The apparatus includes a specimen chamber in which a specimen table is located, a plasma formation chamber separated from the specimen chamber, a microwave introducing window at one end of the plasma chamber and a plasma extracting orifice at the other end of the plasma chamber facing the specimen chamber. A magnetic circuit at the periphery of the plasma formation chamber forms a magnetic flux density necessary to produce ECR. The divergent magnetic field reduces in intensity in a direction towards the specimen chamber. The plasma formation chamber can have a shape and dimensions suitable for forming a microwave cavity resonator and gases can be supplied to the plasma formation and specimen chambers.

Of particular concern to the present invention, PECVD and ECR CVD processes (also referred to hereinafter as plasma CVD processes) may be used for growing silicon dioxide on substrate surfaces, particularly over semiconductor wafers where the silicon dioxide usually forms a dielectric layer between adjacent metallization layers. The use of aluminum in the metallization layers greatly limits the maximum temperature to which the wafers can be exposed. Plasma CVD allows lower processing temperatures to be used compared to thermal CVD processes.

Plasma CVD processes for growing silicon dioxide are very useful because of their low temperature characteristics. However, there are certain

limitations on growing silicon dioxide by plasma CVD compared to thermal CVD silicon dioxide deposition. Plasma CVD silicon dioxide films are usually less stable than thermal CVD silicon dioxide films, i.e., the atoms in the plasma CVD films will be present in higher free energy configurations than in the thermal CVD films. Such "high stress" films are undesirable since they can cause wafer deformation, i.e., bowing, which adversely affects subsequent processing steps. Also, plasma CVD silicon dioxide films will typically have higher levels of incorporated hydrogen (as hydroxyl) when hydrogen is present in one of the reactant gases compared to thermal CVD films. Higher hydrogen incorporation is undesirable because it is associated with poor film properties and a high dielectric constant.

The level of stress in plasma CVD films can be reduced by RF sputtering which appears to preferentially remove those atoms in the film that are bound in higher free energy states (least strongly bound) and thus contribute to film stress. Such sputtering, however, cannot be adjusted independently of other process parameters to reduce stress to optimally low levels. Moreover, such sputtering has no significant effect on lowering hydrogen incorporation into the silicon dioxide films.

PECVD processes for growing silicon nitride using feed mixtures containing nitrogen trifluoride (NF_3), silane (SiH_4), and nitrogen gas (N_2) are described in publications such as Chorng-Ping et al. (1988), Journal of Vacuum Science and Technology B, Vol. 6, No. 2, pp. 524-532; Flamm et al. (1987), Solid

State Technology, March, pp. 43-44; Fujita et al. (1985), J. Appl. Phys. 57:426-432; Livengood and Hess (1987) Appl. Phys. Lett. 50:560-562; Livengood et al. (1988), J. Appl. Phys. 63:2651-2659; Chang et al. 5 (1987), J. Appl. Phys. 62:1406 et seq.; and Pai et al. (1990), J. Appl. Phys. 68:2442-2449. These processes generally result in the deposition of fluorinated silicon nitride.

Processes for periodically plasma etching a 10 growing PECVD film have been proposed. The etching step(s) would be performed only after the PECVD process has been stopped and the deposition reactants removed.

U.S. Patent No. 4,401,054 ("Matsuo") discloses an ECR CVD apparatus wherein gaseous material and 15 microwave power are introduced into a plasma formation chamber to generate plasma by microwave discharge through electron cyclotron resonance. Matsuo discloses that the apparatus can be used to form a thin film of Si, Si_3N_4 , SiO_2 , MoSi_2 and WSi_2 .

20 Matsuo discloses several ECR CVD processes for forming thin film. For instance, a Si_3N_4 film can be formed by introducing N_2 into the plasma formation chamber and SiH_4 into the specimen chamber. Instead of introducing only N_2 into the plasma formation chamber, 25 Ar can be introduced to form a Si film or a mixture of Ar + 20% N_2 can be introduced to form a Si_3N_4 film. A SiO_2 film can be formed by introducing O_2 into the plasma formation chamber and SiH_4 into the specimen chamber. Phosphorus-silicate glass ("PSG") films can 30 be formed by introducing O_2 into the plasma formation chamber and SiH_4 + PH_3 into the specimen chamber.

Molybdenum silicide (MoSi_2) films can be formed by introducing MoF_6 into the plasma formation chamber and SiH_4 into the specimen chamber. Matsuo also discloses that the ECR CVD apparatus can also be used for plasma etching by introducing gases such as CF_4 .

U.S. Patent No. 4,481,229 ("Suzuki") discloses a method of growing a silicon-including film by using a halogenide silicon gas. Suzuki discloses that when Si-N or a-Si films are formed in conventional CVD apparatus of the D.C. glow discharge type and RF discharge type, hydrogen is contained in the deposited film. However, to avoid hydrogen in the Si including film, halogenide silicon gas such as SiF_4 , SiCl_4 , SiFCl_3 , SiF_3Cl and SiBr_4 can be used as the silicon-supplying gas instead of SiH_4 . A problem with using such halogenide silicon gases is that the desired film is not deposited. Instead, a silicon wafer used as the substrate is etched. One reason for this is that the operating gas pressure is higher than 10^{-2} Torr, the electron temperature of discharge is low (about 4 eV) and the bonding energy of the halogenide gas Si-F ($Q=115$ kcal/mol) or Si-Cl ($Q=67.8$ kcal/mol) is greater than that of Si-H ($Q=53.7$ kcal/mol). Thus, the halogenide silicon gas is not fully decomposed. In addition, in the case of the RF discharge type apparatus, the film deposited on the surface is sputtered or decomposed by the incident ions thus preventing the film from growing.

Suzuki discloses that a hydrogen-free silicon-including film can be grown by ECR CVD using a halogenide silicon gas alone or in a gas mixture. Suzuki's apparatus includes a round waveguide, a

discharge tube, a solenoid coil for forming a magnetic field in a vacuum chamber which includes the inside of the discharge tube and a specimen chamber, a permanent magnet outside the specimen chamber but below a
5 substrate table in the specimen chamber aligned with the discharge tube, and gas inlets in the substrate chamber. The discharge tube is made of an insulator (such as quartz or alumina) capable of transmitting the microwave and the strength of the magnetic field in the
10 discharge tube decreases in a direction towards the substrate surface. The permanent magnet gathers the lines of magnetic force together at the substrate surface.

Suzuki forms a Si_3N_4 hydrogen-free film by
15 introducing non-hydrogen containing gases such as SiF_4 and N_2 in a ratio of 1:1 into the vacuum chamber. Suzuki discloses that NF_3 can be used as the nitrogen-containing gas. To produce an a-Si film, SiF_4 is used as the discharge gas. Suzuki discloses that a silicon-
20 containing film can also be grown using SiCl_4 , SiF_nCl_m ($n + m = 4$) or $\text{Si}_x\text{F}_y\text{Cl}_z$ ($x \geq 2$).

U.S. Patent No. 4,973,381 ("Palmer") discloses a method of plasma etching wherein excited gas contacts the surface to be etched and excited electrons in the
25 gas molecules tend to remix with molecules of the surface material to form volatile compounds which are then removed by vacuum pumps. Palmer discloses that the active species of the gas can be a halogenated gas such as CCl_4 or a mixture of CCl_4 and O_2 . Other active
30 species include O, OH, F, CO, CF, Cl, F_2 , CCl, CF_2 , Cl_2 and Br. Buffer gases which change the distance the

reactant travels in Palmer's apparatus prior to recombination include helium, argon and nitrogen.

There is a need in the art for plasma CVD processes for growing films such as silicon dioxide over semiconductors and other substrate surfaces to provide layers having reduced intrinsic stress and reduced levels of incorporated hydrogen in the form of hydroxyl.

SUMMARY OF THE INVENTION

10 The invention provides a process for growing a SiO_x film having reduced intrinsic stress by providing a substrate in a reaction zone, simultaneously introducing different first and second gases into the reaction zone, growing a film comprising atoms in
15 ordered and disordered crystallographic states on the substrate by decomposing the first gas and reacting the decomposed gas on a surface of the substrate, and
20 reducing intrinsic stress in the film by removing atoms from the film in the disordered crystallographic state. The disordered atoms are removed by reacting a chemically active species of the second gas with the disordered atoms and the film growing and intrinsic stress reducing steps are performed simultaneously.

25 The invention also provides a process for growing a SiO_x film having reduced hydrogen content by providing a substrate in a reaction zone, simultaneously introducing different first and second gases into the reaction zone, at least one of the first and second gases containing hydrogen, growing a film on
30 the substrate by decomposing the first gas and reacting

the decomposed gas with the substrate, the growing film including hydrogen atoms therein, and removing hydrogen atoms from the film by reacting the growing film with a chemically active species in the second gas, the film growing and hydrogen removing steps being performed simultaneously.

The process can include introducing a third gas into the reaction zone, the third gas being decomposed along with the first gas during the film growing step.

10 The third gas is different from the first and second gases and the film comprises a compound of elements removed from the first and third gases. The process is particularly useful in growing a silicon dioxide film on a semiconductor wafer. To grow silicon dioxide, the

15 first gas can comprise a silicon containing gas such as SiH₄, Si₂H₆, SiF₄, Si₂F₆, SiCl₄, SiH₂Cl₂ and organosilanes, the second gas can comprise a halogen gas such as NF₃, HF, SF₆, CF₄, C₂F₆, C₂Cl₃F₃, SF₆, Br₂, Cl₂, P₂ and I₂ and the third gas can comprise O₂, N₂O and CO₂. The SiO_x film can also be produced directly

20 from the first gas if the first gas comprises an organosilane.

The film growing, intrinsic stress reducing and hydrogen reducing steps can be performed in a vacuum environment such as a reaction chamber of a plasma enhanced chemical vapor deposition apparatus. The

25 intrinsic stress reducing step can reduce the magnitude of the intrinsic stress in the film to below 200 MPa, preferably below 150 MPa and more preferably below 100

30 MPa. The hydrogen reducing step can reduce hydrogen in the form of hydroxyl from the film to below 3 at. %, preferably below 2 at. % and more preferably below 1.5

> 200 MPa = ? 2×10^9 dynes/cm²
 > 150 MPa =
 > 100 MPa = 10^9 dynes/cm²

at. 4. The intrinsic stress reducing step and/or the
hydrogen removing step preferably etches the growing
film at a rate of 20 to 90% or 20 to 60% of a rate at
which the film grows during the film growing step. The
etching rate can be adjusted by controlling the ratios
of the gases introduced into the reaction zone.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

The present invention provides an improved
process for growing films such as SiO_x films over
substrate surfaces, particularly for the plasma-
enhanced CVD (PECVD) and electron cyclotron resonance
(ECR) CVD of silicon dioxide on semiconductor wafers
during device fabrication procedures. The improvement
comprises the combination of plasma-phase chemical etch
reaction(s) concurrently with the plasma-phase chemical
vapor deposition reaction(s) occurring at or near the
substrate surface. It has been found that the chemical
etching of certain high free energy, unstable atoms
during the deposition process significantly reduces
intrinsic stress within the deposited film.
Additionally, it has been found that the use of halogen
etchants in the deposition process reduces the hydrogen
content within the deposited films. Thus, silicon
dioxide films are produced which are superior to those
obtained without concurrent chemical etching.

Chemical vapor deposition processes are well
known in the art. See, for example, Elliot, Integrated
Circuit Fabrication Technology, McGraw-Hill Book
Company, New York, 1982, the disclosure of which is
hereby incorporated by reference. Conventional
thermal CVD processes rely on heat-induced

decomposition of reactant gases to promote the reaction rate of the gases at the substrate surface to produce a desired film. In PECVD, disassociation of the reactant gases can be enhanced by the application of radio frequency (RF) energy to a reaction zone proximate the substrate surface, creating a plasma of highly-reactive ionic species. See, Chapman, Glow Discharge Processes, John Wiley and Sons, New York 1980, the disclosure of which is hereby incorporated by reference.

10 Recently, a variation of PECVD referred to as electron cyclotron resonance (ECR) CVD has been developed wherein dissociation of the reactant gases into a plasma of ions is achieved in a separate plasma chamber by exposure to microwave energy, typically at
15 2.45 GHz, in the presence of a magnetic field. Electrons are constrained to move in orbits in the magnetic field. The electrons then drift in the direction of the diverging magnetic field which extends toward the substrate. The extraction of the electrons
20 from the plasma chamber via the magnetic field imposes an electrical potential gradient which then draws the ions out of the plasma chamber after the electrons. In the vicinity of the substrate surface, the RF discharge creates a large potential gradient over a
25 short (with respect to the scale of the machine) distance which accelerates the ions perpendicularly towards the substrate. Thus, the ions are only directly affected, to any substantial degree, by the electrostatic fields that exist in the plasma, not by
30 the magnetic or RF fields. By varying the RF power applied to the substrate, the magnitude of the potential gradient and thus the arrival energy of the ions at the substrate surface, can be varied.

ECR CVD has numerous advantages including independent (non-coupled) control over the plasma generation energy and the RF bias energy. Additionally, the shape of the plasma can be modified in a desirable manner by the magnetic field.

The methods of the present invention are particularly advantageous when applied to PECVD and ECR CVD processes. The use of simultaneous etching could also be incorporated in conventional thermal CVD processes as well.

CVD reactors suitable for use in the present invention may be conventional and can include a support surface for one or more substrates, typically semiconductor wafers, and a means for cooling the substrate(s) to below a desired temperature, typically below about 450°C, more typically in the range from about 400°C to about 250°C. The cooling means may comprise fluid cooling channels formed in the body of the support surface for the wafer. PECVD reactors further include a radio frequency discharge means, typically a pair of electrodes, capable of producing a glow discharge which converts the gaseous reactants and the vapor phase etchants into corresponding free radical species. ECR CVD reactors include an electron cyclotron resonance source for producing the desired reactive plasma species and may include an RF biasing means for accelerating the reactive species towards the wafer.

According to one aspect of the present invention, SiO_x can be formed by the vapor phase (usually plasma

5 phase) reaction of a silicon reactant such as silane (SiH_4), disilane (Si_2H_6), silicon tetrafluoride (SiF_4), silicon hexafluoride (Si_2F_6), silicon tetrachloride (SiCl_4), silyldichloride (SiH_2Cl_2), and commercially available organosilanes such as TEOS (tetraethoxy-silane), TOMCATS (2,4,6,8 tetramethylcyclotetra-siloxane), DABS (diacetoxyditertiarybutoxysilane) and the like.

TEOS

10 The oxygen reactant can be selected from gases such as molecular oxygen (O_2), nitrous oxide (N_2O), carbon dioxide (CO_2), and the like. The flow rate of the reactants into the reactor will depend primarily on reactor size, deposition rate, number of wafers, and the like. The ratio of silicon reactant to oxygen
15 reactant will typically be in the range from about 5:1 (oxygen:silicon on an atomic basis) to about 2:1, more typically being in the range from about 4:1 to about 3:1. The silicon and oxygen reactants can be introduced as separate gases or as part of the same gas
20 (e.g. organosilane gas).


For PECVD and ECR CVD, the pressure within the reaction zone of the reactor should preferably be maintained substantially below atmospheric, typically being below about 1 Torr, usually below about 100
25 mTorr, and advantageously being in the range from about 0.5 to 100 mTorr. The pressure can be maintained using conventional rotary and turbomolecular or diffusion pumps in combination with conventional pressure control instrumentation.

30 In the case of PECVD, radio frequency discharge will be provided in order to induce a glow discharge to

maintain the reactant and etchant species in their free radical forms. The radio frequency can typically be applied using internal opposed electrodes operated at a frequency of about 13.56 MHz. The power level may
5 vary from about 0 Watts·cm² to about 2.5 Watts·cm², typically being in the range from about 1 Watt·cm² to about 2 Watts·cm².

In the case of ECR CVD, microwave energy is introduced into the plasma chamber where the conditions
10 for electron cyclotron resonance are met. Typical ECR conditions are a microwave frequency of 2.45 GHz and a magnetic field strength of 875 Gauss. The microwave power levels can typically be in the range from about 0.2 kW to about 5 kW, usually being in the range from
15 about 1 kW to about 3 kW.

As described thus far, process conditions are similar to those employed in conventional PECVD and ECR CVD silicon dioxide deposition. In the present invention, however, the conditions are modified to
20 selectively induce chemical plasma etching of relatively high energy silicon dioxide molecules which if not removed, would increase the intrinsic stress of the deposited film.

 As used herein, intrinsic stress is defined as stress inherent in the deposited film, in contrast to film stress which results from differential thermal expansion between the film and substrate. Intrinsic stress is conveniently measured in units of MPa and can be determined by equipment which measures bowing of the
25
30 wafer caused by stress. Such measurement techniques are well known and the measuring equipment is available

from commercial suppliers, such as Flexus, Sunnyvale, California.

Chemical etching according to the present invention is carried out concurrently with deposition by introducing a vapor phase chemical etchant to the mixture of reaction gases so that an etching mechanism will occur simultaneously with the silicon dioxide deposition mechanism within the reaction zone. The etchant can be a conventional chemical etchant of the type used for plasma etching of silicon dioxide. The etchant can be further selected so that it will not significantly introduce impurities into the film and will be otherwise compatible with the process characteristics. Suitable etchants for the process of the present invention include halogen etchants, such as nitrogen trifluoride (NF_3), HF , SF_6 , CF_4 , C_2F_6 , $\text{C}_2\text{Cl}_3\text{F}_3$, SF_6 , and the like, as well as pure halogens, such as bromine (Br_2), chlorine (Cl_2), fluorine (F_2), iodine (I_2), and the like. The use of NF_3 is preferred. The use of pure halogens, however, is not preferred since they expose the deposition equipment to corrosion. Also, sulfur-based and carbon-based etchants, such as SF_6 , CF_4 , C_2F_6 , and the like, are not generally preferred as they can result in undesirably high levels of impurities such as residual carbon or sulfur in the deposited film.

The etchants will be introduced to the reaction zone (or optionally to the plasma chamber in the case of ECR CVD) in an amount sufficient to achieve an appropriate balance between the deposition rate and the etch rate. Typically, higher amounts of etchant results in a lower deposition rate of the SiO_x film.

5 Suitable etch rates can vary from about 20% to about 90% of the deposition rate. For example, the etch rate can be from about 20% to about 60% of the deposition rate. The atomic ratio of fluorine to silicon introduced to the reaction zone can be between about 1:10 to 4:1 (F:Si), usually being between about 1:10 to 2:1 (F:Si).

10 Use of the halogen etchants according to the present invention has been found to reduce the amount of hydrogen in the form of hydroxyl (OH) which is formed within the deposited silicon dioxide layer. In conventional processes, the amount of hydrogen present as hydroxyl will often exceed 10%, typically being above about 5%. Using the method of the present
15 invention with halogen etchants, the amount of hydrogen in the deposited silicon dioxide layer can be reduced to below about 3 atomic %, frequently being below about 2 atomic %, and often being below about 1.5 atomic %. The amount of hydrogen incorporated in the silicon
20 dioxide film may be measured by conventional methods and equipment, such as those which employ infrared spectroscopy, such as the Bio-Rad FTS-7 analyzer.

25 Although the invention has been described with reference to the foregoing embodiments, various changes and modifications may be made thereto which fall within the scope of the appended claims.

What Is Claimed Is:

1 1. A process for growing a SiO_x film having
2 reduced intrinsic stress, comprising the steps of:
3 providing a substrate in a reaction zone;
4 simultaneously introducing first and second
5 gases into the reaction zone, the first gas being
6 different from the second gas;
7 growing a film on the substrate by
8 decomposing the first gas and reacting the decomposed
9 gas on a surface of the substrate, the growing film
10 comprising atoms in ordered and disordered
11 crystallographic states; and
12 reducing intrinsic stress in the film by
13 removing atoms from the growing film in the disordered
14 crystallographic state, the disordered atoms being
15 removed by reacting a chemically active species of the
16 second gas with the growing film, the film growing and
17 intrinsic stress reducing steps being performed
18 simultaneously.

1 2. The process of claim 1, wherein the first
2 gas comprises an organosilane gas and the second gas
3 comprises a halogen containing gas.

1 3. The process of claim 1, wherein the first
2 gas comprises a silicon containing gas and the second
3 gas comprises a halogen containing gas, the process
4 further comprising a step of introducing a third gas
5 into the reaction zone during the film growing step,
6 the third gas comprising an oxygen containing gas and
7 the film consisting essentially of silicon dioxide.

1 4. The process of claim 1, wherein the first
2 gas comprises a gas selected from the group consisting

3 of SiH₄, Si₂H₆, SiF₄, Si₂F₆, SiCl₄, SiH₂Cl₂ and
4 organosilanes.

1 5. The process of claim 3, wherein the third
2 gas comprises a gas selected from the group consisting
3 of O₂, N₂O and CO₂.

1 6. The process of claim 1, wherein the second
2 gas comprises a halogen gas selected from the group
3 consisting of NF₃, HF, SF₆, CF₄, C₂F₆, C₂Cl₃F₃, SF₆,
4 Br₂, Cl₂, F₂ and I₂.

1 7. The process of claim 1, wherein the film
2 growing and intrinsic stress reducing steps are
3 performed in a vacuum environment.

1 8. The process of claim 7, wherein the vacuum
2 is below 1 Torr and the substrate is maintained at a
3 temperature below about 450°C.

1 9. The process of claim 1, wherein the film
2 growing and intrinsic stress reducing steps are
3 carried out in a reaction chamber of a plasma enhanced
4 chemical vapor deposition apparatus.

1 10. The process of claim 1, wherein the
2 intrinsic stress reducing step reduces a magnitude of
3 intrinsic stress in the film to no greater than a range
4 of 100 to 200 MPa.

1 11. The process of claim 1, wherein the second
2 gas comprises a fluorine containing etchant.

12. The process of claim 11, wherein the fluorine-containing etchant is introduced to the reaction zone at an atomic ratio of F:Si of between 1:10 to 4:1.

1 13. The process of claim 1, wherein the film is
2 deposited on a silicon wafer.

1 14. The process of claim 13, wherein the silicon
2 wafer includes an exposed metallization layer and the
3 SiO_x film is applied as a dielectric layer on the
4 silicon wafer.

1 15. The process of claim 1, wherein at least one
2 of the first and second gases contains hydrogen and the
3 intrinsic stress reducing step results in reducing
4 hydrogen content in the film.

1 16. The process of claim 15, wherein the
2 intrinsic stress reducing step reduces hydrogen in the
3 form of hydroxyl in the film to no greater than a range
4 of 1.5 to 3 at. %.

1 17. The process of claim 1, wherein the
2 intrinsic stress reducing step etches the film at a
3 rate of 20 to 90% of a rate at which the film is grown
4 during the film growing step.

1 18. A process for growing a SiO_x film having
2 reduced hydrogen content, comprising the steps of:
3 providing a substrate in a reaction zone;
4 simultaneously introducing first and second
5 gases into the reaction zone, the first gas being

6 different than the second gas and at least one of the
7 first and second gases containing hydrogen;
8 growing a film on the substrate by
9 decomposing the first gas and reacting the decomposed
10 gas with the substrate, the film including hydrogen
11 atoms therein; and
12 removing the hydrogen atoms from the film by
13 reacting the growing film with a chemically active
14 species in the second gas, the film growing and
15 hydrogen removing steps being performed simultaneously.

1 19. The process of claim 18, wherein the first
2 gas comprises an organosilane gas and the second gas
3 comprises a halogen containing gas.

1 20. The process of claim 18, wherein the first
2 gas comprises a silicon containing gas and the second
3 gas comprises a halogen containing gas, the process
4 further comprising a step of introducing a third gas
5 into the reaction zone during the film growing step,
6 the third gas comprising an oxygen containing gas and
7 the film consisting essentially of silicon dioxide.

1 21. The process of claim 18, wherein the first
2 gas comprises a gas selected from the group consisting
3 of SiH_4 , Si_2H_6 , SiF_4 , Si_2F_6 , SiCl_4 , SiH_2Cl_2 and
4 organosilanes.

1 22. The process of claim 20, wherein the third
2 gas comprises a gas selected from the group consisting
3 of O_2 , N_2O and CO_2 .

1 23. The process of claim 18, wherein the second
2 gas comprises a halogen gas selected from the group

3 consisting of NF_3 , HF , SF_6 , CF_4 , C_2F_6 , $\text{C}_2\text{Cl}_3\text{F}_3$, SF_6 ,
4 Br_2 , Cl_2 , F_2 and I_2 .

1 24. The process of claim 18, wherein the film
2 growing and hydrogen removing steps are performed in a
3 vacuum environment.

1 25. The process of claim 24, wherein the vacuum
2 is below 1 Torr and the substrate is maintained at a
3 temperature below about 450°C .

1 26. The process of claim 18, wherein the film
2 growing and hydrogen removing steps are carried out in
3 a reaction chamber of a plasma enhanced chemical vapor
4 deposition apparatus.

1 27. The process of claim 18, wherein the
2 hydrogen removing step reduces a magnitude of intrinsic
3 stress in the film to no greater than a range of 100 to
4 200 MPa.

1 28. The process of claim 18, wherein the second
2 gas comprises a fluorine-containing etchant.

1 29. The process of claim 28, wherein the
2 fluorine-containing etchant is introduced to the
3 reaction zone at an atomic ratio of F:Si of 1:10 to
4 4:1.

1 30. The process of claim 18, wherein the film is
2 deposited on a silicon wafer.

1 31. The process of claim 30, wherein the silicon
2 wafer includes an exposed metallization layer and the

3 SiO_x film is applied as a dielectric layer on the
4 silicon wafer.

1 32. The process of claim 18, wherein the
2 hydrogen reducing step reduces hydrogen in the form of
3 hydroxyl in the film to no greater than a range of 1.5
4 to 3 at. %.

1 33. The process of claim 18, wherein the
2 hydrogen reducing step etches the film at a rate of 20
3 to 90% of a deposition rate of the film during the film
4 growing step.

A. CLASSIFICATION OF SUBJECT MATTER

IPC(S) : C23C 16/00, 16/18, 16/40

US CL : 427/38, 39, 255.3; 156/643, 646; 437/228

According to International Patent Classification (IPC) or to both national classification and IPC.

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 427/38, 39, 255.3; 156/643, 646; 437/228

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS

search terms: silicon dioxide, silicon oxide, silica organosilanes, silanes, SiH₄, oxygen, carbon dioxide, nitrous oxide
halogen**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US.A, 4,894,352 (Lane et al) 16 January 1990, see col. 1, lines 41-50.	1-12, 14-15, 18-29 and 31
Y	US.A, 3,655,438 (Sterling et al) 11 April 1972, see col. 1, lines 41-52.	13, 16-17, 30 and 32-33
Y	US.A, 4,614,666 (Lindenfelser) 30 September 1986 see col. 1, line 47-col. 2, line 3.	13, 16-17, 30 and 32-33
Y,P	US.A, 5,023,206 (Freeman) 11 June 1991, see col. 2, lines 18-40.	1-12, 14-15, 18-29 and 31

☐ Further documents are listed in the continuation of Box C.☐ See patent family annex.

* Special categories of cited documents:

A document defining the general state of the art which is not considered to be part of particular relevance

E earlier document published on or after the international filing date

L document which may throw doubt on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

Z document member of the same patent family

Date of the actual completion of the international search

16 AUGUST 1992

Date of mailing of the international search report

29 SEP 1992

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